FDA Approves New Pfizer and Moderna XFG mRNA COVID Bioweapon Injections

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BREAKING: FDA APPROVES
NEW PFIZER AND MODERNA XFG mRNA COVID SHOTS BASED ON DATA FROM JUST 8–10 MICE
XFG-SPECIFIC DATA: 8–10 MICE
APPROVED
FDA U.S. FOOD & DRUG ADMINISTRATION
CENTER FOR BIOLOGICS EVALUATION AND RESEARCH
APPLICATION FOR LICENSURE OF A BIOLOGICS PRODUCT
APPROVED
Pfizer XFG COVID-19 mRNA vaccine
For Intramuscular Use
moderna XFG COVID-19 mRNA vaccine
For Intramuscular Use
NO HUMAN TRIALS OF THE NEW FORMULATIONS BEFORE APPROVAL
REACH OF THE APPROVALS:
PFIZER: 65+ AND HIGH-RISK AGES 5–64
MODERNA: 65+ AND HIGH-RISK AGES 6 MONTHS–64
APPROVED FOR HIGH-RISK INFANTS AS YOUNG AS 6 MONTHS

Dylan Eleven | Truth11.com

With millions dead and injured from the Covid bioweapon 'vaccines'; Instead of banning them, the FDA approves more deadly injections. This shows a clear signal that they are still engaged in the genocide program. At what point will these people be stopped and held accountable?

Nicolas Hulscher, MPH | Focal Points | Courageous Discourse

FDA Approves New Pfizer and Moderna XFG mRNA COVID Shots Based on Data From Just 8–10 Mice.

The new formulations were approved before their human safety and immunogenicity studies even began.

The FDA has **approved Pfizer** and **Moderna's** new 2026–2027 XFG-adapted mRNA COVID-19 vaccines even though the XFG-specific evidence **publicly presented to regulators** was preclinical animal data involving just 8–10 mice per group.

The reach of these dangerous approvals is sweeping: Pfizer can now be given to every American age 65 and older and to "high-risk" individuals as young as 5, while Moderna extends all the way down to "high-risk" infants just 6 months old.

The human safety and immunogenicity studies of the new formulations had not even begun when FDA granted approval.

Pfizer's new COMIRNATY formulation was approved on **August 27, 2026**, while its prospective human study of the 2026–2027 formula is scheduled to begin **August 31—four days later**.

Moderna's new SPIKEVAX formulation was also approved on **August 27**, but its Phase 3b/4 human study of the 2026–2027 SPIKEVAX and MNEXSPIKE formulations is not scheduled to begin until **November 16—nearly three months after approval**.

FDA's own 2026 vaccine-selection materials distinguish between human data from earlier/current COVID vaccines and animal immunogenicity data for the new candidate vaccines.

Pfizer's FDA presentation shows the XFG candidate was tested in **10 mice per group**.

Moderna's FDA presentation shows its key XFG experiment used **8 mice per group**.

The approval letters reveal another troubling development: FDA also released **both manufacturers from previously required placebo-controlled postmarketing studies**, citing operational and feasibility problems. Pfizer's abandoned study would have investigated circulating spike antigen and post-vaccination symptoms, while Moderna's would have evaluated safety of variant-containing formulations.

Rather than requiring new human data before approval, FDA relied in part on human safety and immunogenicity data from previous COVID-19 vaccine formulations and extrapolated that evidence to the newly updated XFG products, while the XFG-specific evidence itself remained preclinical.

This reliance on older-formulation data is especially concerning given the **millions of deaths, injuries, and disabilities**, along with demonstrated **cardiotoxicity**, **neurotoxicity**, **carcinogenicity**, **body-wide biodistribution**, and **years-long persistence**.

This is extremely worrisome and needs to be reversed immediately.

Source: https://www.thefocalpoints.com/p/breaking-fda-approves-new-pfizer

Original Article: Truth Warrior Suggested

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